

FORM PTO-1449 (REV.7-80)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTY. DOCKET NO. 480140.442C1	APPLICATION NO. 09/765,105
INFORMATION DISCLOSURE STATEMENT (Use several sheets if necessary)		APPLICANTS		Donald S. Karanewsky et al.	
		FILING DATE		GROUP ART UNIT	
		January 16, 2001		1653	

U.S. PATENT DOCUMENTS

EXAMINER DRAFTED	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
DL	AA 6,235,899	05/22/01	Bouchet et al.	540	500	
	AB					
	AC					
	AD					

FOREIGN PATENT DOCUMENTS

	DOCUMENT NUMBER	DATE	COUNTRY	TRANSLATION	
				YES	NO
DL	AE WO 97/22619	06/26/97	WIPO		
	AF WO 98/10778	03/19/98	WIPO		
	AG WO 99/03852	01/28/99	WIPO (+ English Translation)	X	
	AH WO 00/01666	01/13/00	WIPO		
	AI WO 00/23421	04/27/00	WIPO		
	AJ WO 01/00658	01/04/01	WIPO		
DL	AK WO 01/51462	07/19/01	WIPO		
DL	AL WO 01/81330	11/01/01	WIPO		

OTHER PRIOR ART (Including Author, Title, Date, Pertinent Pages, Etc.)

DL	AM	Chapman K., "Synthesis of a Potent, Reversible Inhibitor of Interleukin-1 β Converting Enzyme," <i>Bioorganic & Medicinal Chemistry Letters</i> 2(6):613-618, 1992.
DL	AN	Cheung et al., "Synthesis of 3-Amino-3-Vinylpropanoic Acid and its Conversion to 4-Amino-5-Hydroxy-4,5-Dihydrofuran-2-one Hydrochloride (HAD), A Cyclic Stabilised Form of Aspartate 1-Semialdehyde Hydrochloride," <i>Tetrahedron</i> 53(46):15807-15812, 1997.
DL	AO	de Lange et al., "Asymmetric 1, 4-Additions to 5-Alkoxy-2(5H)-Furanones Enantioselective Synthesis and Absolute Configuration Determination of β -Amino- γ -Butyrolactones and Amino Diols," <i>Tetrahedron</i> 45(21):6799-6818, 1989.

EXAMINER	David Luckay	DATE CONSIDERED
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* EXAMINER: Initial if reference considered, whether or not criteria is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant(s).

FORM PTO-1449 (REV.7-80)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTY. DOCKET NO. 480140.442CI	APPLICATION NO. 09/765,105
OCT 27 2003 PAPER FILED INFORMATION DISCLOSURE STATEMENT (Use several sheets if necessary)		APPLICANTS		Donald S. Karanewsky et al.	
		FILING DATE		GROUP ART UNIT 1653	

U.S. PATENT DOCUMENTS

EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
	BA						
	BB						
	BC						
	BD						

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	TRANSLATION	
					YES	NO
	BE					
	BF					
	BG					

OTHER PRIOR ART (Including Author, Title, Date, Pertinent Pages, Etc.)

<i>DR</i>	BH	Faber et al., "Catalytic Kinetic Resolution of 5-Alkoxy-2(5H)-Furanones," <i>Tetrahedron</i> 50(16):4775-4794, 1994.
<i>DR</i>	BI	Feringa et al., "Asymmetric Synthesis of 2-Amino-1, 4-Diols," <i>Tetrahedron Letters</i> 29(11):1303-1306, 1988.
<i>DR</i>	BJ	Feringa et al., "1, 4-Additions of Amines to 5-Methoxyfuran-2(5H)-One; An Efficient Synthesis of Amino Diols," <i>Heterocycles</i> 27(5):1197-1205, 1988.
<i>DR</i>	BK	Furuichi et al., "Common Synthetic Strategy for Optically Active Cyclic Terpenoids having a 1,1,5-Trimethyl- <i>Trans</i> -Decalin Nucleus: Syntheses of (+)-Acuminolide, (-)-Spongianolide A, and (+)-Scalarenedial," <i>Tetrahedron</i> 57, pp. 8425-8442, 2001.
<i>DR</i>	BL	Gonzalez et al., "Pseudoesters and Derivatives. Part38. 1,3-Dipolar Cycloadditions of Aryl Azides and an Aziridine, <i>Via</i> Azomethine Ylide, to 2(5H)-Furanones Substituted at the 5-Position by Methoxy and Sulfur Bearing Groups," <i>Heterocycles</i> 52(1):237-251, 2000.
<i>DR</i>	BM	Leblanc et al., "Sar in the Alkoxy Lactone Series: The Discovery of DFP, A Potent and Orally Active Cox-2 Inhibitor," <i>Bioorganic & Medicinal Chemistry Letters</i> 9, pp. 2207-2212, 1999.
<i>DR</i>	BN	Lubben et al., "Asymmetric Synthesis of β -Lactams via Amine Additions to 5(R)-Menthyl-2(5H)-Furanone," <i>Tetrahedron: Asymmetry</i> 2(8):775-778, 1991.

EXAMINER	<i>David Luton</i>	DATE CONSIDERED
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